

AMENDMENTS TO THE CLAIMS

1. (**Currently amended**) A recombinant nucleotide sequence which codes upon expression a bifunctional hybrid active-site serine β -lactamase protein, wherein the β -lactamase protein is a class A, C or D β -lactamase protein that bears at least one heterologous sequence; wherein the β -lactamase protein bears the at least one heterologous sequence in a region forming a juncture between alpha helix 8 and alpha helix 9 of said active-site serine β -lactamase in a region located between two neighboring alpha helices of the β -lactamase sequence, wherein the region is selected from the group consisting of:

- a) a region forming a juncture between alpha helix 8 and alpha helix 9 of TEM-1 β -lactamase; and
- b) a region forming a juncture between the alpha helices of said class A, C or D β -lactamase, said alpha helices corresponding to the alpha helix 8 and alpha helix 9 of the TEM-1 β -lactamase, and

wherein the hybrid protein has two functions, wherein, in said bifunctional hybrid protein, the first function is associated with the β -lactamase portion and the second function is associated with the at least one heterologous sequence having a biological function which is different from the first function.

2.-5. (Canceled)

6. (**Previously presented**) The recombinant nucleotide sequence according to Claim 1, wherein the β -lactamase moiety is a class A β -lactamase, wherein said β -lactamase class A protein bears the at least one heterologous sequence in the region forming a juncture between alpha helix 8 and alpha helix 9.

7. (**Previously presented**) The recombinant nucleotide sequence according to claim 1, wherein the region forming a juncture between alpha helix 8 and alpha helix 9 is selected from the group consisting of:

- a) amino acid sequence Thr195 to Leu199 of the TEM-1 β -lactamase; and
- b) an amino acid sequence in a β -lactamase other than TEM-1 β -lactamase corresponding to the amino acid sequence Thr195 to Leu199 in TEM-1 β -lactamase.

8.-11. (Canceled)

12. **(Currently amended)** A recombinant nucleotide sequence which codes upon expression a bifunctional hybrid class A β -lactamase class-A protein, wherein the class A β -lactamase class-A protein bears at least one heterologous sequence in a region located between two neighboring alpha helices of the β -lactamase sequence, wherein the region is selected from the group consisting of:

a) the a region forming a juncture between alpha helix 8 and alpha helix 9 of the TEM-1 β -lactamase; and

b) the a region forming a juncture between the alpha helices of said a-homologous class A β -lactamase-class-A, said alpha helices corresponding to the alpha helix 8 and alpha helix 9 of the TEM-1 β -lactamase,

wherein the hybrid protein has a first function and a second function, wherein the first function is associated with the β -lactamase portion and is selected from the group consisting of:

c) hydrolyzing β -lactams (β -lactamase activity); and

d) binding covalently and in a stable manner to substances selected from the group consisting of β -lactams, derivatives of β -lactams, and inhibitors of β -lactams; and wherein the second function is associated with the at least one heterologous sequence having a biological function which is different from the first function.

13.-15. **(Canceled)**

16. **(Previously presented)** The recombinant nucleotide sequence according to Claim 1, wherein the at least one heterologous sequence has a length of 11 or more amino acid residues.

17. **(Previously presented)** The recombinant nucleotide sequence according to Claim 1, wherein the at least one heterologous sequence has a length of 18 or more amino acid residues.

18. **(Previously presented)** The recombinant nucleotide sequence according to Claim 1, wherein the at least one heterologous sequence has a length of 25 or more amino acid residues.

19. **(Previously presented)** The recombinant nucleotide sequence according to Claim 1, wherein the at least one heterologous sequence has a length of 50 or more amino acid residues.

20. **(Previously presented)** The recombinant nucleotide sequence according to Claim 1, wherein the at least one heterologous sequence has a length of 100 or more amino acid residues.

21. **(Currently amended)** The recombinant nucleotide sequence according to Claim 1, wherein the nucleotide sequence coding for the β -lactamase sequence encodes is selected from the group consisting of:

- a) nucleotide sequence coding for the β -lactamase TEM-1 (SEQ ID NO: 1)
- b) nucleotide sequence coding for the β -lactamase BlaP (SEQ ID NO: 2);
- c) nucleotide sequence coding for the β -lactamase BlaL (SEQ ID NO: 3);
- d) nucleotide sequence coding for the β -lactamase AmpC (SEQ ID NO: 39); and
- e) nucleotide sequence coding for the β -lactamase BlaR-CTD (SEQ ID NO: 41);
- f) a recombinant sequence of one or more of a) to e); and
- g) nucleotide sequences which hybridize under stringent conditions to the nucleotide sequences of any one of a) to f).

22. **(Currently amended)** The recombinant nucleotide sequence according to Claim 1, wherein the at least one heterologous sequence is ~~related to a function~~ selected from the group consisting of: ~~being~~ an epitope, ~~being~~ a specific binding partner for antibodies, being a sequence that is specifically recognized and bound by antibodies, a sequence having a binding affinity to earth alkali and metal ions, a sequence having enzymatic activity, ~~being~~ a toxin, (StA heat-stable enterotoxin of *E. coli*), ~~bearing~~ a glycosylation site, ~~bearing~~ a glycosylated peptide, ~~being~~ a specific binding partner for any polypeptide or any ligand, and a sequence having a binding affinity to dsDNA, and ssDNA or RNA ~~(having a binding affinity to nucleotide and polynucleotide)~~.

23. **(Currently amended)** The recombinant nucleotide sequence according to Claim 1, wherein the at least one nucleic acid sequence encoding the at least one heterologous sequence is selected from the group consisting of: STa (heat stable enterotoxin of *Escherichia coli*, SEQ ID NO: 21); encodes protein A of *Staphylococcus aureus* with two Fc Binding domains, (SEQ ID NO: 23 and 25); ~~protein G of *Streptococcus pyogenes*, (SEQ ID NO: 27 and 29)~~, a linear antigenic determinant of the hemagglutinin of the Influenza virus (SEQ ID NO: 31), a fragment of human phospholipase type 11 (hPLA2) (SEQ ID NO: 33), and LPS binding amino acid sequence (SEQ ID NO: 35); and nucleotide sequences which hybridize under stringent conditions to said nucleotide sequences.

24.-53. **(Canceled)**